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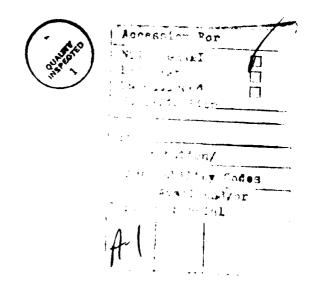
This study of the cofiring of used automotive oil in Navy boilers was sponsored by the Naval Facilities Engineering Command (NAVFAC 032P) under program element 63721N, Task Area Y0817 SL. It examines the used oil regulations and environmental quality requirements involved with used automotive oil cofiring. The results of the air dispersion modeling of a boiler cofiring a used automotive oil are also included.

This report has been reviewed by J. L. Brumfield, Head, Chemical Systems Branch and H. O. Oliver, Head, Survivability Division.

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INTRODUCTION

The U.S. Navy is concerned with finding alternate disposal methods for used oils generated at naval facilities. The cofiring of used automotive oil in Navy boilers is an alternative to the current disposal methods for this used oil. Cofiring provides ultimate disposal of this waste along with a savings on boiler fuel. However, recent Federal regulations limit the burning of used oil for energy recovery; and future regulations may include used oil as a hazardous waste. With regards to the cofiring of used automative oil, this study was primarily concerned with the lead emissions from boilers burning this used oil.

This report (1) examines how these regulations will affect the cofiring of used automotive oil in Navy boilers, (2) demonstrates how air dispersion modeling can be used to show what impact cofiring used automotive oil will have on the ambient air quality (directly affecting the ambient air quality is the lead concentration of the used oil burned in the boilers), and (3) discusses the issue of whether or not the Navy should consider adopting a lower specification for the lead concentration in used oil fuel than is presently required in the regulations. For background information, refer to the Initiation Decision Report on Thermal Destruction of Hazardous Wastes in Navy Boilers and Incinerators. 1

CURRENT REGULATIONS ON USED OIL BURNING

On 29 November 1985, the Environmental Protection Agency (EPA) issued new regulations under the Resource Conservation and Recovery Act (RCRA) covering the burning of used oil and hazardous waste for energy recovery (40 CFR Part 266—Standards for the Management of Specific Hazardous Wastes and Specific Types of Hazardous Waste Management Facilities²). These regulations distinguish between a used oil and a hazardous waste in setting the restrictions in burning for energy recovery. Also included is a used oil specification that limits the types of boilers that can burn used oils that fail the specification.

Various used oils can be classified as a characteristic hazardous waste under 40 CFR 261, Subpart C--Characteristics of Hazardous Waste, which lists two of the characteristics as Ignitability and EP Toxicity. This is important, since used automotive oil may meet the EP Toxicity characteristic for lead and be considered a hazardous waste. However, under the new regulations, a used oil that is also a characteristic hazardous waste is regulated under Subpart E--Used Oil Burned for Energy Recovery and not under Subpart D--Hazardous Waste Burned for Energy Recovery.

In Subpart E of the regulations, a used oil is defined as "any oil that has been refined from crude oil, used, and as a result of such use, is contaminated by physical or chemical impurities." Any fuel produced by blending, processing,

or other treatment of a used oil is a used oil fuel. Used oil fuel that cannot meet the specifications given in Table 1 is an off-specification used oil fuel. Lead is the constituent that may cause untreated used automotive oil to fail the specifications. Blending of the off-specification used oil with virgin fuel oil will usually be the easiest means of treating it to meet the specifications.

TABLE 1. USED OIL FUEL SPECIFICATIONS

Constituent/Property	Allowable Level
Arsenic	5 ppm maximum
Cadmium	2 ppm maximum
Chromium	10 ppm maximum
Lead	100 ppm maximum
Flash Point	100 °F maximum
Total Halogens	4000 ppm maximum

Off-specification used oil fuel is regulated as to what type of devices can use it as a fuel. They are limited to industrial furnaces, industrial boilers, utility boilers, and used oil-fired space heaters as defined in the regulations. Industrial boilers are "boilers located on the site of a facility engaged in a manufacturing process where substances are transformed into new products." Utility boilers are "boilers used to produce electric power, steam, or heated or cooled air or other gases or fluids for sale." It is unknown if Navy boilers will be considered in either of these two categories. Used oil fuel that meets the specification can be burned in any boiler regardless of function. Off-specification used oil fuel is also subject to record keeping and analysis requirements that are different from those for used oil that meets the specifications.

The following is a summary of the analysis and record-keeping requirements most applicable to Navy facilities wishing to cofire used automotive oil. A Navy facility blending an off-specification used oil fuel (used automotive oil) to meet specifications must obtain analysis or other information to prove that the resulting used oil fuel does meet the specifications. These records must be kept for 3 yrs. The facility must also notify the EPA or state, if authorized, of its used oil management activities. This will be required even if the facility previously notified the EPA of hazardous waste management activities and obtained a U.S. EPA Identification Number. The burning of off-specification used oil fuel requires the burner to notify the EPA as to the location and general description of its used oil management activities.

A Navy facility selling used automotive oil that is off-specification is not required to notify the EPA of its used oil management activities unless the used oil is sold directly to a person who burns it for energy recovery. If sold to such a person, the facility must notify the EPA of its used oil management activities, send the proper invoices with each shipment, and obtain the proper notices from the burner. The invoices and notices must be kept for a period of 3 yrs. If the facility sells used oil claiming to meet the specifications, it must obtain analysis to prove the used oil meets the specifications. These

records must be kept for 3 yrs. An operations log book must also be kept that details each shipment of used oil. These must also be kept for 3 yrs. The facility must notify the EPA or state, if authorized, of its used oil management activities. Notification is required because the facility will be the first to claim that the used oil meets the specifications.

These regulations became effective 9 December 1985. The only exception is the lead specification that became effective 29 May 1986. Besides the Federal regulations, many states and localities have regulations governing used oil burning. These regulations must also be met.

FUTURE REGULATIONS ON USED OIL BURNING

Future regulations call for stricter control over recycled used oil and also the listing of used oil as a hazardous waste. Phase I regulations on the blending and burning of used oil and hazardous waste are the regulations discussed previously. Phase II regulations are due to be proposed sometime in 1986. They may contain emission standards for burners of used oil. The lead specification for off-specification used oil fuel will be reviewed and may be lowered.

The proposed Recycled Used Oil Standards will regulate recycled oil.³ These standards will apply to generators and transporters of recycled oil along with the owners and operators of used oil recycling facilities. Used oil that is either burned for energy recovery or used to produce a fuel is considered a recycled oil. A fuel produced by blending used automotive oil and virgin fuel oil is subject to these regulations unless it can meet certain specifications that are identical to those in the current regulations. Other proposed regulations call for the listing of used oil as a hazardous waste. This regulation will list used oil as a hazardous waste but will leave the management of used oil recycling to the above regulations.

IMPACT OF REGULATIONS ON COFIRING

Used automotive oil is a waste generated by many Navy facilities. In the past, several facilities have cofired this waste.⁴ To determine the current level of used oil cofiring, NAVFAC Engineering Field Divisions and several Navy facilities were contacted regarding this practice. A summary of these results is presented in Appendix A. No facilities were found to be cofiring used automotive oil. Thus, the regulations will have no immediate effect on current cofiring practices.

The impact of the regulations on the disposal of used automotive oil through contractors will depend on how the regulations affect the contractor. Some contractors may no longer be able to accept the used oil and the facility will be forced to find another contractor or an alternate disposal method. The Navy facility selling the used oil is subject to certain requirements under the regulations as previously described.

The impact of the regulations on future plans to cofire used automotive oil can be minimized by meeting the used oil fuel specifications. This will limit the regulations that need to be complied with to cofire the used oil. As the lead content of used automotive oil drops, it will be come easier to meet the current lead specification and any lower specification that may be established. The impact of the yet-to-be-proposed Phase II regulations cannot be determined until they are established.

LEAD CONCENTRATION IN USED AUTOMOTIVE OIL

An accurate lead concentration in used automotive oil is difficult to obtain due to the phasedown of lead in gasoline. Lead in gasoline is the sole source of lead in automotive oil. In July 1985, the EPA lowered the maximum lead content in gasoline from 1.1 gr/gal to 0.5 gr/gal. In January 1986, it was lowered to 0.1 gr/gal. The EPA may even consider a total ban on lead in gasoline. Subsequently, the lead content in used automotive oil is dropping. It is unsure if the lead content will drop below 100 ppm. If it does and the used oil meets all other specifications, it will no longer be an off-specification used oil fuel. Table 2 shows the lead concentration of various groups of samples collected between 1981 and 1983. The current lead concentration will be lower than these figures.

TABLE 2. LEAD CONCENTRATIONS IN SAMPLES

			Concentration Units (ppm)					
	Number of	Number/Percent Containing				r in ntile gory	-	
Type of Oil	Samples	Lead	Mean	Median	75	90	Range	
Used Automotive	96	94/97	1603	900	1600	2300	1 - 21,700	
Direct From Generator*	40	39/97	2574	1470	2210	3300	5 - 21,900	
From Waste Oil Collectors or Processors**	56	55/98	915	730	1110	1417	1 - 11,000	
From Diesel Engines	5	4/80	29	13			<5 - 78	
From Casoline Engines**	* 87	87/100	2232	390			9 - 21,676	

^{*}Known to be only used automotive oil

^{**}Collector/processor claimed the oil was only used automotive oil; however, the exact nature of the oil is unknown

^{***}No distinctions between oil samples from leaded or unleaded gasoline; from the general population of motor vehicles, not Navy vehicles

LEAD CONCENTRATION IN FUEL OIL RECLAIMED (FOR)

FOR is a product of Navy reclamation operations. It is reclaimed from large volumes of waste oil and used as a boiler fuel. Included in the waste oil is used lubricating oil. This creates some concern about the lead concentration in FOR, which was found to be less than 60 ppm. 6 The FOR specification, MIL-F-24951, does not include a lead concentration as a requirement.

AIR DISPERSION MODELING

A dispersion model is a mathematical representation of the meteorological transport and turbulent diffusion processes that occur in the atmosphere. It will predict the pollutant concentration due to emissions from various sources for a set of locations, called receptors, for various average time intervals. Facilities wishing to cofire used oil can estimate the effect it will have on ambient air quality through dispersion modeling. This may be required by local or state air permitting agencies to show compliance with their regulations. Air dispersion modeling for which the results are to be used for air permitting requirements should be done by experts in the field of dispersion modeling.

There are many off-the-shelf air dispersion models available. A set of air dispersion models developed by the EPA is available on the User's Network for Applied Modeling of Air Pollution (UNAMAP) system. Besides the UNAMAP system, there are other air dispersion models available for use. An air dispersion model will give valid results only when applied to the specific conditions for which it was developed. Some models can be applied to emissions from sources located in urban areas, while others work best for rural areas. Some other factors to be considered in choosing a model are number and type of emission sources, receptor distances from source, and terrain conditions.

For valid results, it is important that the data inputted to the model be as accurate as possible. The data required for most models include source data, receptor data and meteorological data. Source data required may be stack height and diameter, stack gas exit velocity and temperature, and pollutant emission rate. Receptor data required involve location of the receptors with respect to the emission source and, on some models, terrain elevation is needed. The meteorological data required by dispersion models usually involve some combination of upper air and surface observations, which are available from the National Climatic Center. The Naval Oceanography Command Detachment (NOCD) is the point of contact for obtaining weather information. It is important that the data be representative of the area to be modeled. NOCD can aid in determining from which upper air and surface observation stations the data should be obtained.

MODELING OF BOILER AT A TYPICAL NAVY FACILITY

The EPA's Single Source (CRSTER) model was used to determine the lead concentration in the ambient air as a result of cofiring used automotive oil in a boiler

located at a typical Navy facility. CRSTER is a point-source model that calculates hourly concentrations and averages these to determine concentration estimates for 3-hr, 8-hr, 24-hr, and annual time increments. It will calculate the concentration at 180 receptors located on a radial grid. The receptors were located every 10 deg on concentric circles at distances of 1, 2, 3, 4, and 5 km from the boiler. This particular version of CRSTER was adapted to be run on a microprocessor.

Boiler information required for the CRSTER model is given in Table 3. The boiler was assumed to be burning a blend of used automotive oil and virgin fuel oil with lead concentrations of 100, 75, and 50 ppm. The fuel blends were burned for 24 hr/day for an entire year.

TABLE 3. BOILER DATA

Boiler Size	165 M-BTU/hr			
Fuel Rate	1000 gal/hr			
Stack Height	44.3 m			
Stack Diameter	2.1 m			
Stack Gas Exit Temperature	470 °K			
Stack Gas Volumetric Flow Rate	66,704 SCFM			
Stack Gas Exit Velocity	8.8 m/s			

An important consideration in the modeling of lead is the percentage of lead in the fuel that ultimately reaches the atmosphere. Some of the lead will remain in the boiler as ash. The percentage of lead leaving the boiler is reported to vary from 3 to 64 percent. Emission control devices will also remove a percentage of the lead from the exhaust gases. Although the boiler used in this study did have emission control devices, many Navy boilers are not equipped with such devices. To model a worst-case scenario, a zero-percent efficiency for the emission control devices was assumed. It was also assumed that 100 percent of the lead in the fuel exited the boiler.

The 10 highest annual mean concentrations were all less than one percent of the National Ambient Air Quality Standard (NAAQS). However, the 10 highest 24-hr concentrations were significantly higher than the 10 highest annual mean concentrations. Table 4 shows a comparison of annual and 24-hr concentrations for fuel lead concentrations of 50 and 100 ppm. Currently, the NAAQS is a 3-mo average of 1.5 ug/m³. The particular CRSTER version used was unable to generate a 3-mo average; therefore, a comparison of these two averages with the NAAQS must be made. The 24-hr concentrations are short-term averages and are not as significant when being compared with a 3-mo average. The annual concentrations agree with the results obtained by others doing similiar air dispersion modeling.

TABLE 4. COMPARISON OF AMBIENT AIR CONCENTRATIONS

100 ppm			50 ppm					
Ann	ua1	24-Hr		Ann	Annual		24-Hr	
Concen- tration	7 NAAQS	Concen- tration	7 NAAQS	Concen- tration	% NAAQS	Concen- tration	% NAAQS	
0.00783	0.52	0.0899	6.0	0.00395	0.26	0.0454	3.0	
0.00710	0.47	0.0782	5.2	0.00358	0.24	0.0395	2.6	
0.00699	0.47	0.0758	5.1	0.00353	0.24	0.0383	2.6	
0.00675	0.45	0.0749	5.0	0.00341	0.23	0.0378	2.5	
0.00664	0.44	0.0670	4.5	0.00335	0.22	0.0338	2.3	
0.00651	0.43	0.0660	4.4	0.00329	0.22	0.0333	2.2	
0.00566	0.38	0.0651	4.3	0.00286	0.19	0.0329	2.2	
0.00564	0.38	0.0634	4.2	0.90285	0.19	0.0326	2.1	
0.00536	0.36	0.0633	4.2	0.00270	0.18	0.0317	2.1	
0.00528	0.35	0.0605	4.0	0.00267	0.18	0.0305	2.0	

The 10 highest 24-, 8-, and 1-hr concentrations were determined. These are significantly higher than the annual concentrations and are maximum short-term concentrations. They are usually of short duration and are not as significant as a long-term average. Appendix B contains complete modelling results.

The volume of used automotive oil disposed of in the modeled boiler is based on the lead concentration of the used oil and fuel oil. A 9-ppm concentration will be assumed for the fuel oil⁹ and a 3300-ppm concentration will be assumed for the used oil. This value was chosen because it represents a value that the lead concentration of 90 percent of the samples of known used automotive oil were below. Based on these figures and a continuous yearly operation of the boiler, 240,000 gal/yr of used automotive oil can be disposed of through boiler cofiring. This is an extremely large volume for one facility to generate and, while there are no figures available, it is likely that most Navy facilities generate a significantly lesser volume. Therefore, depending on boiler size, many facilities will be able to cofire a blended fuel with a lead concentration below 100 ppm.

The use of computer air dispersion modeling can be a useful tool in determining what effect used automotive oil cofiring will have on the ambient air quality. Which Navy agency will handle any dispersion modeling is not determined at this time.

AVAILABLE GUIDANCE FOR THE BURNING OF USED OIL

There are several reports available to Navy facilities to provide guidance on the burning of used oil. The Period Polymer Series Seri

as a Supplemental Boiler Fuel issued by the Naval Civil Engineering Laboratory (NCEL) are two such reports. Guidance is provided in the areas of segregation, storage, treatment, fuel analysis, blending, burner modifications, maintenance, and record-keeping. Several Technical Notes also issued by NCEL discuss the results of waste oil burning in Navy boilers. 10.11 These works provide much of the guidance needed to burn used oil in Navy boilers; however, revisions will be needed to account for the new used oil regulations.

RECOMMENDATIONS

A study to be prepared for the Defense Environmental Leadership Project Office will evaluate the economic impact of various disposal options for used automotive oil; cofiring is one of these options. In conjunction with economic guidance, there will be a need for technical guidance. Reports exist that provide technical guidance for the use of used oil as a supplemental fuel in Navy boilers; however, there will be a need for this information to be updated wich respect to the newest used oil regulations. It is recommended that a technical guidance manual be developed for the use of waste oil as a supplemental boiler fuel. Topics discussed should include storage and transportation, sampling, analysis, pretreatment, blending, emission control, permitting, record-keeping, and boiler modification and maintenance. The development of this manual should be delayed until the used oil regulations are finalized.

The 100-ppm lead specification required for used oil fuel to be cofired in Navy boilers is sufficient to meet the NAAQS for lead. The Navy should not consider adopting a lower lead specification than required in the current regulations. However, depending on the quantity of used automotive oil generated and boiler fuel usage rate, many facilities will be able to cofire a fuel blend with a lower lead concentration. This is highly recommended, since it will result in less impact on air quality and will allow easier compliance with any new lead specifications in future regulations.

Since used automotive oil may be a constituent of FOR, it is recommended that lead be included as a test parameter in the specifications for FOR. The concentration level can be that of the lead concentration in the used oil specification.

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APPENDIX A

RESULTS OF USED AUTOMOTIVE OIL BURNING SURVEY

Recently-proposed federal regulations (40 CFR, Part 266, Subparts D and E) regulate the burning of hazardous wastes and used oils. These regulations will affect the cofiring of used automotive oil in Navy boilers. For this reason, it is necessary to determine the current level of used automotive oil cofiring being practiced at Navy facilities. Environmental personnel at NAVFAC Engineering Field Divisions were contacted regarding this practice in their respective divisions. A summary of the results is presented below:

Engineering Field Division	Point of Contact	Comment
Northern	George Wiese AV 443-6280 (215) 897-6280	No facilities currently cofire used automotive cil; suggested contact-ing Naval Avionics Center in Indianapolis, Indiana
Southern	Craig Campbell AV 794-5510 (803) 743-5510	No facilities currently cofire used automotive oil; suggested contact-ing Naval Shipyard in Charleston, South Carolina
Western	John Gordon AV 439-8666 (206) 476-8666	No facilities currently cofire used automotive oil; it is sent to an oil reclamation facility where it is mixed with other oils and sold
Western (San Diego Area)	D.C. Fischer AV 958-5461 (619) 696-5461	One facility may be burning used oilMiramar Naval Air Station
Atlantic	Paul Parker AV 564-9559 (804) 444-9559	A solid refuse boiler at Public Works Center in Norfolk, Virginia burns used oil; used automotive oil goes into Fuel Oil Reclaimed (FOR)
Chesapeake	Steve Buckley AV 288-3760 (202) 433-3760	No facilities currently cofire used automotive oil; it is disposed of through contractors

Several waste oil reports prepared for the Navy have given examples of Navy facilities that cofired waste automotive oil either directly or blended with other waste oils. These facilities, plus the several mentioned by NAVFAC personnel, were contacted to determine their current used automotive oil disposal practices. Both the past and current practice of each facility is described below in as much detail as was available:

NAVAL AIR STATION, FALLON, NEVADA Ronald Nygren (702) 423-5161

Past Practice

Approximately 200 gal of waste oil were generated and burned each month. It was collected in drums at motor pool service areas and transported to the boiler plant and strained through a fine wire screen prior to introduction into primary fuel storage. The waste oil consisted primarily of used lube oil with some hydraulic oil and, rarely, small amounts of grease. A 200-hp boiler was operated for only 7 to 8 mo out of the year at 40 to 55 gal of fuel/hr. The fuel oil normally used was No. 6 fuel oil; No. 2 fuel oil was used as a backup fuel. The normal fuel was stored in a 12,000-gal tank and pumped through a filter into three burners. No special modifications to boiler equipment or operational changes were required to burn this waste oil and no adverse effects from its use were noted.

Current Practice

Waste oil is no longer burned in the boiler. The boiler was converted to natural gas as its main fuel. Possible contamination of the waste oil led local environmental agencies to begin questioning stack emissions and requesting daily reports. Waste oil is currently being disposed of on a single-time contract basis.

NAVAL AIR STATION WHIDBEY, ISLAND, WASHINGTON Ken Buehn AV 820-2811 or 3348

Past Practice

The station collected and burned approximately 60,000 gal of waste oil and contaminated fuel/yr from two different type areas on the base. The waste oil burned consisted of a wide range of solvents, phenols, JP fuel, hydraulic fluids, and used lube oils. Enlisted personnel collected fuel, solvents, and used lube oil from the flight line in mobile bowsers (700- to 1000-gal capacity) and transported it to the boiler plant, where it was stored in a 25,000-gal underground tank. There was no treatment of this oil. Secondly, Public Works collected used lube oil from garages, service areas, and other locations throughout the base. This oil was placed in a 5000-gal tank to settle. The oil that rose was skimmed off the top and placed in a second tank (3000-gal capacity) to settle, and water was drained off the bottom periodically. After settling, this oil was transported to the underground tank at the boiler plant. The waste oil was burned in one of three 44M-BTU/hr boilers that used natural gas as the primary fuel and No. 2 as the backup fuel. One of the three burners was set aside to burn the waste oil. No modifications to the burner itself were required. The waste oil was pumped from the underground holding tank through a fine strainer and into the burner. While the waste oil was being burned, adjustments to the natural gas burners were made to compensate for fluctuations of the waste oil burner. Increased boiler cleaning activities were observed. The boiler utilizing waste oil experiencing increased ash accumulation and required cleaning twice a year. Boilers burning regular oil had to be cleaned only once every year or two.

Current Practice

Waste oil is no longer being burned. With JP fuel now being recycled, it is no longer available for fuel. Waste oil is now being disposed of through the Defense Property Disposal Office (DPDO).

PUBLIC WORKS CENTER, NORFOLK, VIRGINIA Bill Thornton (804) 444-4853

Past Practice

FOR has been burned in boilers at this facility since 1979. It is obtained from Craney Island and was the primary supplemental fuel to the normal No. 6 fuel oil used. A secondary source of fuel was used lube oil and hydraulic fluid, which never amounted to more than 10,000 gal/yr. The waste oil was collected from the motor pool and service areas and was transported to the boiler plant. Here it was directly blended into existing No. 6 fuel oil tankage and was burned along with the regular fuel.

Current Practice

All waste oil is sent to Craney Island and then sold back to the facility as FOR; this includes waste automotive oil. FOR is still being burned in the boilers.

NAVAL AIR STATION, MIRAMAR, CALIFORNIA Ms. Severance 959-3321

Current Practice

At one time, used oil was burned; however, FOR is currently being burned. Burning of waste oil was stopped due to the operational problems it presented. Problems with the state of California over whether it was a hazardous waste also led to its discontinuance. Used automotive oil is currently being sold.

NAVAL WEAPONS SUPPORT CENTER, CRANE, INDIANA Dave Fry AV 482-1521

Current Practice

Waste automotive oil was burned in several boilers at this facility. The boilers have since been converted to natural gas and do not cofire the waste oil. The oil is collected and then sampled by the Environmental Protection group before it is disposed of.

NAVAL AVIONICS CENTER, INDIANAPOLIS, INDIANA Dick Schatz (317) 353-3525

Past Practice

This facility burned approximately 5000 gal of waste oil/yr in one of three 17,000-1b/hr boilers used for steam production. The major source of the waste oil was machine shop cutting oil. Metal filings present in the oil were the major problem and were centrifuged prior to storing and burning. Used lube oil was the secondary source of waste oil and was collected from 60 vehicles in the transportation department and from a collection center provided for the individual to recycle waste oil. The boilers were designed to burn natural gas as primary fuel and have the capability of using No. 5 fuel oil as a backup. Storage of No. 5 fuel oil was maintained in six 1500-gal tanks, one of which was set aside to receive waste oil. Clean No. 5 fuel oil and waste oil were mixed at about a 50-50 ratio in this tank prior to burning.

Current Practice

The waste oil is no longer cofired in the boilers, since problems in proper operation of the boiler were encountered when burning this waste oil. Problems of increased smoke emissions and clogged filters due to dirt and sediment suspended in the waste oil were sometimes experienced. The waste oil is now disposed of through a contractor.

APPENDIX B

RESULTS OF COMPUTER MODELING AT A TYPICAL NAVY FACILITY

TABLE B-1. ANNUAL MEAN CONCENTRATION (ug/ m^3) AT EACH RECEPTOR (Pb = 100 ppm)

Direction		Distance (km)									
(deg)	1.0	2.0	3.0	4.0	5.0						
10	0.00469	0.00528	0.00472	0.00402	0.00345						
20	0.00263	0.00326	0.00316	0.00282	0.00249						
30	0.00190	0.00277	0.00273	0.00241	0.00210						
40	0.00141	0.00189	0.00172	0.00147	0.00126						
50	0.00108	0.00146	0.00137	0.00119	0.00104						
60	0.00107	0.00140	0.00127	0.00108	0.000934						
70	0.000694	0.000847	0.000745	0.000626	0.000537						
80	0.000716	0.000917	0.000879	0.000772	0.000679						
90	0.00115	0.00140	0.00128	0.00112	0.000949						
100	0.00107	0.00133	0.00121	0.00106	0.000958						
110	0.00124	0.00181	0.00183	0.00169	0.00157						
120	0.00226	0.00415	0.00464	0.00448	0.00422						
130	0.00277	0.00465	0.00507	0.00484	0.00449						
140	0.00311	0.00483	0.00492	0.00448	0.00399						
150	0.00326	0.00527	0.00523	0.00464	0.00405						
160	0.00226	0.00369	0.00353	0.00304	0.00259						
170	0.00177	0.00287	0.00268	0.00227	0.00191						
180	0.00191	0.00300	0.00275	0.00231	0.00193						
190	0.000978	0.00133	0.00199	0.000999	0.000841						
200	0.00106	0.00149	0.00138	0.00117	0.000998						
210	0.00153	0.00206	0.00184	0.00153	0.00128						
220	0.00150	0.00163	0.00137	0.00113	0.000949						
230	0.00176	0.00187	0.00156	0.00128	0.00107						
240	0.00185	0.00194	0.00165	0.00136	0.00115						
250	0.00147	0.00144	0.00117	0.000939	0.000770						
260	0.00122	0.00115	0.000983	0.000827	0.000707						
270	0.00118	0.00118	0.00108	0.000951	0.000837						
280	0.000941	0.000877	0.000724	0.000589	0.000491						
290	0.00101	0.00103	0.000870	0.000711	0.000589						
300	0.00160	0.00177	0.00154	0.00128	0.00108						
310	0.00219	0.00210	0.00168	0.00133	0.00108						
320	0.00351	0.00302	0.00230	0.00177	0.00141						
330	0.00536	0.00456	0.00351	0.00275	0.00222						
340	0.00651	0.00566	0.00434	0.00337	0.00271						
350	0.00699	0.00664	0.00518	0.00406	0.00329						
360	0.00710	0.00783	0.00675	0.00564	0.00480						

TABLE B-2. HIGHEST 24-HR CONCENTRATION (ug/ m^3) AT EACH RECEPTOR (Pb = 100 ppm)

Direction			Distance (km)	_	
(deg)	1.0	2.0	3.0	4.0	5.0
10	0.0444 (221)	0.0493 (21)	0.0407 (99)	0.0366 (99)	0.0315 (99)
20	0.0460 (201)	0.0468 (166)	0.0456 (166)	0.0379 (166)	0.0309 (166)
30	0.0317 (139)	0.0322 (100)	0.0296 (100)	0.0249 (46)	0.0215 (46)
40	0.0372 (206)	0.0378 (45)	0.0291 (62)	0.0248 (62)	0.0213 (62)
50	0.0394 (176)	0.0366 (46)	0.0291 (46)	0.0228 (46)	0.0183 (46)
60	0.0307 (105)	0.0269 (70)	0.0246 (128)	0.0225 (128)	0.0196 (128)
70	0.0293 (105)	0.0228 (105)	0.0218 (274)	0.0199 (274)	0,0176 (274)
80	0.0273 (273)	0.0175 (273)	0.0171 (195)	0.0170 (195)	0.0148 (195)
90	0.0543 (140)	0.0298 (140)	0.0283 (28)	0.0255 (28)	0.0226 (28)
100	0.0287 (140)	0.0207 (55)	0.0192 (55)	0.0161 (55)	0.0134 (55)
110	0.0254 (240)	0.0226 (56)	0.0207 (56)	0.0194 (129)	0.0189 (129)
120	0.0411 (115)	0.0359 (115)	0.0357 (312)	0.0357 (312)	0.0337 (312)
130	0.0441 (213)	0.0394 (5)	0.0455 (340)	0.0505 (340)	0.0490 (340)
140	0.0550 (130)	0.0488 (57)	0.0425 (57)	0.0338 (57)	0.0283 (280)
150	0.0565 (123)	0.0457 (123)	0.0435 (279)	0.0398 (279)	0.0349 (279)
160	0.0349 (101)	0.0507 (109)	0.0432 (109)	0.0343 (109)	0.0296 (345)
170	0.0274 (145)	0.0531 (18)	0.0528 (18)	0.0449 (18)	0.0374 (18)
180	0.0356 (254)	0.0406 (244)	0.0362 (38)	0.0315 (150)	0.0275 (150)
190	0.0206 (259)	0.0261 (259)	0.0198 (259)	0.0150 (38)	0.0120 (38)
200	0.0242 (120)	0.0429 (344)	0.0467 (344)	0.0413 (344)	0.0353 (344)
210	0.0441 (255)	0.0386 (235)	0.0401 (235)	0.0355 (235)	0.0305 (235)
220	0.0382 (262)	0.0400 (235)	0.0398 (235)	0.0343 (223)	0,0290 (235)
230	0.0670 (262)	0.0417 (262)	0.0299 (262)	0.0230 (262)	0.0223 (63)
240	0.0596 (148)	0.0288 (151)	0.0245 (151)	0.0210 (125)	0.0179 (143)
250	0.0401 (102)	0.0305 (102)	0.0198 (102)	0.0159 (342)	0.0133 (342)
260	0.0347 (98)	0.0243 (165)	0.0256 (165)	0.0222 (165)	0.0187 (165)
270	0.0292 (246)	0.0153 (127)	0.0172 (127)	0.0173 (71)	0,0166 (71)
280	0.0297 (110)	0.0204 (110)	0.0196 (169)	0.0171 (169)	0.0144 (169)
290	0.0175 (267)	0.0206 (287)	0.0133 (287)	0.0113 (236)	0.0097 (187)
300	0.0304 (230)	0.0397 (188)	0.0383 (188)	0.0322 (188)	0.0266 (188)
310	0.0314 (238)	0.0226 (156)	0.0190 (156)	0.0172 (187)	0.0156 (187)
320	0.0426 (275)	0.0323 (271)	0.0323 (271)	0.0274 (271)	0.0228 (271)
330	0.0783 (214)	0.0653 (214)	0.0447 (214)	0.0318 (214)	0.0238 (214)
340	0.0758 (160)	0.0444 (160)	0.0321 (251)	0.0259 (184)	0.0213 (183)
350	0.0561 (160)	0.0429 (78)	0.0298 (78)	0.0231 (26)	0.0196 (26)
360	0.0900 (221)	0.0618 (221)	0.0400 (221)	0.0352 (67)	0.0322 (315)

NOTE: Number in parenthesis is day of the year in which concentration occurred.

TABLE B-3. HIGHEST 8-HR CONCENTRATIONS (ug/m³) AT EACH RECEPTOR (Pb = 100 ppm)

Direction					Distanc	e (km)				
(deg)	1.	,0	2.	0	3,	,0	4.	.0	5.	.0
10	0.112	•		•	-	•		(99,3)		•
20	0.138							(166,3)		
30	0.0886	(45,2)								
40	0.108	(45,2)						(45,2)		
50		(176,2)								
60		(105, 2)								
70	0.0870	(105,2)	0.0649	(105,2)	0.0654	(274,1)	0.0596	(274,1)	0.0527	(274,1)
80	0.0820	(273,2)	0.0510	(273,2)	0.0435	(195,3)	0.0424	(195,3)	0.0358	(195,3)
90	0.163	(140,2)	0.0893	(140,2)	0.0478	(140,2)	0.0430	(28,1)	0.0411	(28,1)
100	0.0860	(140,2)	0.0484	(140,2)	0.0346	(55,2)	0.0275	(173,3)	0.0244	(173,3)
110	0.0760	(240,2)	0.0669	(350,2)	0.0479	(350,2)	0.0401	(256,1)	0.0413	(256,1)
120	0.0981	(115,2)	0.0776	(226,1)	0.0841	(226,1)	0.0776	(226,1)	0.0696	(226,1)
130	0.116	(213,2)	0.0942	(5,2)	0.133	(340,2)	0.143	(340,2)	0.134	(340,2)
140	0.159	(130,2)	0.104	(130,2)	0.0840	(319,3)	0.0732	(319,3)	0.0619	(319,3)
150	0.152	(123,2)	0.0912	(123,2)	0.0742	(102,1)	0.0675	(102,1)	0.0585	(102,1)
160	0.789							(345,1)		
170	0.0743	(145,2)								
180	0.107							(150,1)		
190		(132 ?)								
200	0.0725	(120,2)								
210	0.132							(260,1)		
220	0.114	(262,2)	0.0881	(236,1)	0.0861	(235,3)	0.0751	(235,3)	0.0644	(235,3)
230	0.201	(262,2)		•		-		(297,2)		•
240	0.168							(151,2)		
250		(102,2)								
260	0.103							(165,1)		
270		(246,2)								
280		(110,2)								
290		(287,2)		•		•		•		•
300		(230,2)		•		•		•		•
310		(238,2)								
320	0.128							(271,1)		
330	0.190							(214,3)		
340	0.220	(167, 2)		,				(26,2)		
350	0.140	•						(188,1)		
360	0.213	(221,2)	0.153	(25,2)	0.105	(25,2)	0.0814	(341,2)	0.0729	(341,2)

NOTE: (A,B) A is the day of the year in which the 8-hr concentration occurred.

B is the 8-hr interval in which the 8-hr concentration occurred.

 $^{1 = 0000 \}text{ to } 0800$

 $^{2 = 0800 \}text{ to } 1600$

 $^{3 = 1600 \}text{ to } 2400$

TABLE B-4. ANNUAL MEAN CONCENTRATION (ug/m³) AT EACH RECEPTOR (Pb = 75 ppm)

Direction					
(deg)	1.0	2.0	3.0	4.0	5.0
10	0.00353	0.00397	0.00355	0.00302	0.00260
20	0.00198	0.00245	0.00333	0.00302	
30	0.00143	0.00243	0.00238	0.00212	0.00187
40	0.00145	0.00203	0.00203	0.00181	0.00158
50	0.000810	0.00110	0.00130	0.00111	0.000945
60	0.000803	0.00110	0.000956	0.000899	0.000784
70	0.000522	0,000637	0.000560	0.000813	0.000703
80	0.000539	0.000690	0.000661	0.000471	0.000404
90	0.000335	0.00105	0.000961	0.000381	0.000511
100	0.000798	0.00100	0.000981	0.000841	0.000748
110	0.000736	0.00136	0.000311	0.000800	0.000721
120	0.000730	0.00138	0.00138	0.00128	0.00118
130	0.00208	0.00313	0.00349		0.00317
140	0.00234	0.00364	0.00370	0.00364	0.00338
150	0.00245	0.00396	0.00370	0.00337 0.00349	0.00301
160	0.00170	0.00378	0.00393		0.00305
170	0.00170	0.00278	0.00202	0.00228	0.00195
180	0.00144	0.00210	0.00202	0.00171	0.00143
190	0.00174	0.00223	0.00207	0.00174	0.00145
200	0.000799	0.00112	0.00104	0.000752	0.000633
210	0.00151	0.00112	0.00104	0.000883	0.000751
220	0.00131	0.00133	0.00139	0.00115	0.000962
230	0.00113	0.00122	0.00103	0.000848	0.000714
240	0.00132	0.00141	0.00118	0.000964	0.000808
250	0.00111	0.00148	0.00124	0.00102	0.000863
260	0.000921	0.00103	0.000739	0.000707	0.000579
270	0.000890	0.000886	0.000739	0.000622	0.000532
280	0.000708	0.000660	0.000545	0.000716	0.000629
290	0.000759	0.000777	0.000545	0.000443	0.000372
300	0.00120	0.00133	0.000855	0.000535	0.000443
310	0.00165	0.00158	0.00115	0.000961	0.000810
320	0.00264	0.00138		0.000999	0.000816
330	0.00204	0.00228	0.00173	0.00133	0.00106
340	0.00403	0.00343	0.00264	0.00207	0.00167
350	0.00490	0.00426	0.00327	0.00254	0.00264
360	0.00534		0.00390	0.00305	0.00248
200	0.00334	0.00589	0.00508	0.00425	0.00361

TABLE B-5. ANNUAL MEAN CONCENTRATION (ug/ m^3) AT EACH RECEPTOR (Pb = 50 ppm)

Direction			Distance (km)		
(deg)	1.0	2.0	3.0	4.0	5.0
10	0.00237	0.00267	0.00238	0.00203	0.00174
20	0.00133	0.00165	0.00160	0.00142	0.00126
30	0.000961	0.00140	0.00138	0.00122	0.00106
40	0.000710	0.000954	0.000870	0.000742	0.000634
50	0.000544	0.000737	0.000690	0.000603	0.000526
60	0.000539	0.000709	0.000641	0.000547	0.000472
70	0.000350	0.000427	0.000376	0.000316	0.000271
80	0.000362	0.000463	0.000444	0.000390	0.000343
90	0.000591	0.000706	0.000644	0.000564	0.000502
100	0.000535	0.000671	0.000611	0.000537	0.000483
110	0.000628	0.000914	0.000924	0.000855	0.000790
120	0.00114	0.00210	0.00234	0.00226	0.00213
130	0.00140	0.00235	0.00256	0.00244	0.00227
140	0.00157	0.00244	0.00249	0.00226	0.00202
150	0.00164	0.00266	0.00264	0.00234	0.00205
160	0.00114	0.00186	0.00178	0.00153	0.00131
170	0.000892	0.00145	0.00135	0.00114	0.000962
180	0.000965	0.00151	0.00139	0.00116	0.000974
190	0.000494	0.000669	0.000601	0.000504	0.000424
200	0.000536	0.000754	0.000697	0.000593	0.000504
210	0.000772	0.00104	0.000929	0.000773	0.000645
220	0.000757	0.000820	0.000692	0.000569	0.000479
230	0.000888	0.000944	0.000780	0.000647	0.000542
240	0.000933	0.000980	0.000831	0.000686	0.000579
250	0.000742	0.000725	0.000591	0.000474	0.000389
260	0.000618	0.000580	0.000496	0.000417	0.000357
270	0.000597	0.000594	0.000545	0.000480	0.000422
280	0.000475	0.000443	0.000365	0.000297	0.000248
290	0.000510	0.000521	0.000439	0.000359	0.000297
300	0.000807	0.000894	0.000775	0.000645	0.000543
310	0.00111	0.00106	0.000846	0.000670	0.000547
320	0.00177	0.00153	0.00116	0.000892	0.000711
330	0.00270	0.00230	0.00177	0.00139	0.00112
340	0.00329	0.00286	0.00219	0.00170	0.00137
350	0.00353	0.00335	0.00261	0.00205	0.00166
360	0.00358	0.00395	0.00341	0.00285	0.00242

TABLE B-6. 10 HIGHEST 24-HR CONCENTRATIONS (Pb = 100 ppm)

Day	24-Hr Concentration (ug/m ³)	Direction (deg)	Distance (km)	7 NAAQS (1.5 ug/m ³)
221	0.0899	0	2.0	6.0
214	0.0782	330	1.0	5.2
160	0.0758	340	1.0	5.1
167	0.0749	340	1.0	5.0
262	0.0670	230	1.0	4.5
158	0.0660	340	1.0	4.4
159	0.0651	340	1.0	4.3
161	0.0634	330	1.0	4.2
138	0.0633	340	1.0	4.2
163	0.0605	330	1.0	4.0

TABLE B-7. 10 HIGHEST 8-HR CONCENTRATIONS (Pb = 100 ppm)

Day	8-Hr Concentration (ug/m ³)	Direction (deg)	Distance (km)	<pre>% NAAQS (1.5 ug/m³)</pre>
167	0.220	340	1.0	14.7
221	0.213	0	2.0	14.2
262	0.201	2 30	1.0	13.4
160	0.193	340	1.0	12.8
158	0.192	340	1.0	12.7
161	0.190	330	1.0	12.7
159	0.184	340	1.0	12.2
161	0.183	340	1.0	12.2
163	0.178	330	1.0	11.9
214	0.171	330	1.0	11.4

TABLE B-8. 10 HIGHEST 1-HR CONCENTRATIONS (Pb = 100 ppm)

Day	1-Hr Concentration (ug/m ³)	Direction (deg)	Distance (km)	7 NAAQS (1.5 ug/m ³)
233	0.354	120	2.0	23.6
26	0.328	340	2.0	21.8
342	0.324	250	2.0	21.5
340	0.313	130	2.0	20.9
26	0.300	0	3.0	20.0
213	0.276	140	1.0	18.4
208	0.275	350	1.0	18.3
215	0.274	330	1.0	18.3
32	0.274	10	2.0	18.3
176	0.274	20	1.0	18.2

TABLE B-9. 10 HIGHEST 24-HR CONCENTRATIONS (Pb = 75 ppm)

Day	24-Hr Concentration (ug/m ³)	Direction (deg)	Distance (km)	7 NAAQS (1.5 ug/m ³)
221	0.0677	0	2.0	4.5
214	0.0589	330	1.0	3.9
100	0.0570	340	1.0	3.8
167	0.0564	340	1.0	3.8
262	0.0504	230	1.0	3.4
158	0.0497	340	1.0	3.3
159	0.0490	340	1.0	3.3
161	0.0477	330	1.0	3.2
138	0.0476	340	1.0	3.2
163	0.0455	330	1.0	3.0

TABLE B-10. 10 HIGHEST 8-HR CONCENTRATIONS (Pb = 75 ppm)

Day	8-Hr Concentration (ug/m ³)	Direction (deg)	Distance (km)	% NAAQS (1.5 ug/m ³)
167	0.166	340	1.0	11.0
221	0.161	0	2.0	10.7
262	0.151	230	1.0	10.1
160	0.145	340	1.0	9.7
158	0.144	340	1.0	9.6
161	0.143	330	1.0	9.5
159	0.139	340	1.0	9.2
161	0.138	340	1.0	9.2
163	0.134	330	1.0	8.9
214	0.128	330	1.0	8.6

TABLE B-11. 10 HIGHEST 1-HR CONCENTRATIONS (Pb = 75 ppm)

Day	1-Hr Concentration (ug/m³)	Direction (deg)	Distance (km)	% NAAQS (1.5 ug/m ³)
==1				<u> </u>
233	0.267	120	2.0	17.8
26	0.246	340	2.0	16.4
342	0.244	250	2.0	16.2
340	0.236	130	2.0	16.2
26	0.225	0	3.0	15.0
213	0.208	140	1.0	13.9
208	0.207	350	1.0	13.8
215	0.206	330	1.0	13.8
32	0.206	10	2.0	13.7
176	0.206	20	1.0	13.7

TABLE B-12. 10 HIGHEST 24-HR CONCENTRATIONS (Pb = 50 ppm)

Day	24-Hr Concentration (ug/m ³)	Direction (deg)	Distance (km)	7 NAAQS (1.5 ug/m ³)
221	0.0454	0	2.0	3.0
214	0.0395	330	1.0	2.6
160	0.0383	340	1.0	2.6
167	0.0378	230	1.0	2.5
262	0.0338	340	1.0	2.3
158	0.0333	340	1.0	2.2
159	0.0329	340	1.0	2.2
161	0.0320	330	1.0	2.1
1 38	0.0319	340	1.0	2.1
163	0.0305	330	1.0	2.0

TABLE B-13. 10 HIGHEST 8-HR CONCENTRATIONS (Pb = 50 ppm)

D	8-Hr Concentration	Direction	Distance	Z NAAQS
Day	(ug/m ³)	(deg)	<u>(km)</u>	(1.5 ug/m^3)
167	0.111	340	1.0	7.4
221	0.108	0	2.0	7.2
262	0.101	2 30	1.0	6.7
160	0.0972	340	1.0	6.5
158	0.0967	340	1.0	6.4
161	0.0958	330	1.0	6.4
159	0.0930	340	1.0	6.2
161	0.0925	340	1.0	6.2
163	0.0899	330	1.0	6.0
214	0.0899	330	1.0	5.7

TABLE B-14. 10 HIGHEST 1-HR CONCENTRATIONS (Pb = 50 ppm)

Day	1-Hr Concentration (ug/m ³)	Direction (deg)	Distance (km)	7 NAAQS (1.5 ug/m ³)
233	0.179	120	2.0	11.9
26	0.165	340	2.0	11.0
342	0.163	250	2.0	10,9
340	0.158	130	2.0	10.5
26	0.151	0	3.0	10.1
213	0.139	140	1.0	9.3
208	0.139	350	1.0	9.3
215	0.139	330	1.0	9.3
32	0.138	10	2.0	9.2
176	0.138	20	1.0	9.2

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